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                 Polymer links for the POLYLINK command completed in REGISTRY
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                 Additional enzyme-catalyzed reactions added to CASREACT
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                 BEILSTEIN on STN workshop to be held August 24 in conjunction
                 with the 228th ACS National Meeting
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         AUG 02
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                 STN User Update to be held August 22 in conjunction with the
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                 Pricing for the Save Answers for SciFinder Wizard within
                 STN Express with Discover! will change September 1, 2004
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         AUG 27
                 BIOCOMMERCE: Changes and enhancements to content coverage
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                 BIOTECHABS/BIOTECHDS: Two new display fields added for legal
         AUG 27
                 status data from INPADOC
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         SEP 01
                 INPADOC: New family current-awareness alert (SDI) available
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                 New pricing for the Save Answers for SciFinder Wizard within
         SEP 01
                 STN Express with Discover!
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         SEP 01
                 New display format, HITSTR, available in WPIDS/WPINDEX/WPIX
NEWS 21
         SEP 14
                 STN Patent Forum to be held October 13, 2004, in Iselin, NJ
              JULY 30 CURRENT WINDOWS VERSION IS V7.01, CURRENT
NEWS EXPRESS
              MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 11 AUGUST 2004
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FILE 'MEDLINE' ENTERED AT 13:59:21 ON 20 SEP 2004

FILE 'BIOSIS' ENTERED AT 13:59:21 ON 20 SEP 2004 Copyright (c) 2004 The Thomson Corporation.

=> s croft-j?/au

466 CROFT-J?/AU

=> s l1 and new (w) zealand

L22 L1 AND NEW (W) ZEALAND

=> d ibib abs 1-2

ANSWER 1 OF 2 CAPLUS COPYRIGHT 2004 ACS on STN L2

ACCESSION NUMBER:

2000:889642 CAPLUS

DOCUMENT NUMBER:

134:21423

TITLE:

SOURCE:

A synergistic composition comprising mussel protein extract and glycosaminoglycan suitable for treatment

of arthritis

INVENTOR(S):

Croft, John Eric

DATE

PATENT ASSIGNEE(S):

MacFarlane Laboratories New Zealand Limited, N. Z. Brit. UK Pat. Appl., 10 pp.

APPLICATION NO.

DATE

CODEN: BAXXDU

Patent

KIND

DOCUMENT TYPE:

LANGUAGE:

English FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.

GB 2347349  Al 20000906 GB 1999-4672  PRIORITY APPLN. INFO.:  A pharmaceutical composition comprising proteins extracted from the New Zealand green-lipped mussel (Perna canaliculus) and one or more glycosaminoglycans, preferable glucosamine or its sulfate, has anti-inflammatory properties. The composition is used in the treatment of arthritis. The combination of the protein extract and the glycosaminoglycan is synergistic with respect to the effect of the same concentration of the individual components. The preferred composition includeds ma homogeneous mixture of a freeze-dried powder containing protein extract and						
PRIORITY APPLN. INFO.:  GB 1999-4672  19990301  AB A pharmaceutical composition comprising proteins extracted from the New  Zealand green-lipped mussel (Perna canaliculus) and one or more glycosaminoglycans, preferable glucosamine or its sulfate, has anti-inflammatory properties. The composition is used in the treatment of arthritis. The combination of the protein extract and the glycosaminoglycan is synergistic with respect to the effect of the same concentration of the individual components. The preferred composition includeds ma homogeneous						
AB A pharmaceutical composition comprising proteins extracted from the New Zealand green-lipped mussel (Perna canaliculus) and one or more glycosaminoglycans, preferable glucosamine or its sulfate, has anti-inflammatory properties. The composition is used in the treatment of arthritis. The combination of the protein extract and the glycosaminoglycan is synergistic with respect to the effect of the same concentration of the individual components. The preferred composition includeds ma homogeneous			20000906	GB 1999-4672	19990301	
AB A pharmaceutical composition comprising proteins extracted from the New Zealand green-lipped mussel (Perna canaliculus) and one or more glycosaminoglycans, preferable glucosamine or its sulfate, has anti-inflammatory properties. The composition is used in the treatment of arthritis. The combination of the protein extract and the glycosaminoglycan is synergistic with respect to the effect of the same concentration of the individual components. The preferred composition includeds ma homogeneous	PRIORITY APPLN. INFO.:			GB 1999-4672	19990301	
glycosaminoglycans, preferable glucosamine or its sulfate, has anti-inflammatory properties. The composition is used in the treatment of arthritis. The combination of the protein extract and the glycosaminoglycan is synergistic with respect to the effect of the same concentration of the individual components. The preferred composition includeds ma homogeneous	AB A pharmaceutical	composit:	ion comprisi	ng proteins extract	ed from the New	
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arthritis. The combination of the protein extract and the glycosaminoglycan is synergistic with respect to the effect of the same concentration of the individual components. The preferred composition includeds ma homogeneous	anti-inflammatory	propert:	ies. The co	mposition is used in	n the treatment of	
is synergistic with respect to the effect of the same concentration of the individual components. The preferred composition includeds ma homogeneous	arthritis. The c	ombinatio	on of the pr	otein extract and t	he glycosaminoglyca	n
individual components. The preferred composition includeds ma homogeneous	is synergistic wi	th respec	ct to the ef	fect of the same co	ncentration of the	
mixture of a freeze-dried powder containing protein extract and	individual compon	ents. Th	ne preferred	composition include	eds ma homogeneous	
J. P.	mixture of a free	ze-dried	powder cont	aining protein extra	act and	

glycosaminoglycan powder. The compns. are capsules or tablets.

ACCESSION NUMBER:

ANSWER 2 OF 2 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on STN

DOCUMENT NUMBER:

1993:438586 BIOSIS PREV199396093211

TITLE:

The independent effects of obesity and body fat

distribution on blood pressure in black adults: The Pitt

County study.

AUTHOR (S): Croft, Janet B.; Strogatz, David S.; Keenan, Nora

L.; James, Sherman A. [Reprint author]; Malarcher, Ann M.;

Garrett, Joanne M.

CORPORATE SOURCE:

Univ. Mich., Dep. Epidemiology, Sch. Public Health, 109

Observatory Street, Ann Arbor, MI 48109, USA

SOURCE:

International Journal of Obesity, (1993) Vol. 17, No. 7,

pp. 391-397.

CODEN: IJOBDP. ISSN: 0307-0565.

DOCUMENT TYPE:

Article

LANGUAGE:

English

ENTRY DATE:

Entered STN: 22 Sep 1993

Last Updated on STN: 22 Sep 1993

The relationship of obesity measures to blood pressure and hypertension prevalence was assessed in a community probability sample of 25-50-year-old black adults (1101 women and 655 men) who were examined in 1988 in Pitt County, North Carolina (USA). Among black women, both body mass index and waist-to-hip ratio had independent relationships with systolic and diastolic blood pressures and hypertension prevalence after controlling for the effects of age, socio-economic status, physical activity, alcohol, and the other obesity measure (P lt 0.05). Body mass index also had independent relationships with blood pressure levels and hypertension prevalence in black men (P lt 0.05), while waist-to-hip ratio was associated with hypertension prevalence (P = 0.05) and diastolic blood pressure (P lt 0.05), but not with systolic blood pressure. relationships of waist-to-hip ratio with blood pressure and hypertension prevalence were considerably reduced in both sex groups after controlling for body mass index. This study presents new evidence that waist-to-hip ratio is related to hypertension and blood pressure level independent of body mass index, in young to middle-aged black adult women and men.

=> s l1 and perna

2 L1 AND PERNA

=> s 13 not 12

1 L3 NOT L2

=> d ibib abs

ANSWER 1 OF 1 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1980:573751 CAPLUS

DOCUMENT NUMBER:

93:173751

TITLE:

Pharmaceutical preparations containing a mollusk

INVENTOR(S):

McFarlane, Stuart John; Croft, John Eric

PATENT ASSIGNEE(S):

SOURCE:

Eur. Pat. Appl., 34 pp.

CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA	TENT NO.			KIND	DATE	APPLICATION NO.	DATE
ΕP	10061			A1	19800416	EP 1979-810098	19790919
EΡ	10061			B1	19830713		
	R: BE,	CH,	DE,	FR,	GB, IT, LU,	NL, SE	
WO	8000661			A1	19800417	WO 1979-EP72	19790920
	W: DE,	SE,	US				
ĎΕ	2953186			${f T}$	19810108	DE 1979-2953186	19790920
CA	1134745			A1	19821102	CA 1979-336104	19790921
$I\Gamma$	58301			A1	19821130	IL 1979-58301	19790921
AU	7951107			A1	19800403	AU 1979-51107	19790924
ΑU	536153			B2	19840419		

ZA 7905039	A	19800924	ZA	1979-5039	19790924
JP 55147223	A2	19801117	JР	1979-122127	19790925
US 4455298	A	19840619	US	1982-376898	19820510
PRIORITY APPLN. INFO.:			NZ	1978-188489	19780925
			·WO	1979-EP72	19790920
			US	1980-194152	19800915

AB The occurrence of gastric ulcers or stomach bleeding from drugs is inhibited by combination with the drugs of a mollusk Perna canaliculus extract (Seatone), which is composed of proteins, carbohydrates and minerals (mineral and amino acid content given). Capsule compns. were given containing the extract and analgesics-inflammation inhibitors such as acetylsalicylic acid [50-78-2], diclofenac Na [15307-79-6], phenylbutazone [50-33-9], or indomethacin [53-86-1].

=> s Perna (w) canaliculus

238 PERNA (W) CANALICULUS L5

=> s 15 and arthritis

24 L5 AND ARTHRITIS 1.6

=>`s 16 and chondroitin

1 L6 AND CHONDROITIN

=> d ibib abs

ANSWER 1 OF 1 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2000:889642 CAPLUS

DOCUMENT NUMBER:

134:21423

TITLE:

A synergistic composition comprising mussel protein

extract and glycosaminoglycan suitable for treatment

of arthritis

INVENTOR (S):

Croft, John Eric

PATENT ASSIGNEE(S):

MacFarlane Laboratories New Zealand Limited, N. Z.

SOURCE:

Brit. UK Pat. Appl., 10 pp. CODEN: BAXXDU

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	<del>-</del>	<b></b>		
GB 2347349	A1	20000906	GB 1999-4672	19990301
PRIORITY APPLN. INFO.:			GB 1999-4672	19990301

AB A pharmaceutical composition comprising proteins extracted from the New Zealand green-lipped mussel (Perna canaliculus) and one or more glycosaminoglycans, preferable glucosamine or its sulfate, has anti-inflammatory properties. The composition is used in the treatment of arthritis. The combination of the protein extract and the glycosaminoglycan is synergistic with respect to the effect of the same concentration of the individual components. The preferred composition includeds ma

homogeneous mixture of a freeze-dried powder containing protein extract and glycosaminoglycan powder. The compns. are capsules or tablets.

=> s shark (w) cartilage

1.8 502 SHARK (W) CARTILAGE

=> s 18 and 16

L9 0 L8 AND L6

=> s 18 and arthritis

L10 14 L8 AND ARTHRITIS

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13 DUPLICATE REMOVE L10 (1 DUPLICATE REMOVED) L11

=> d ibib abs 1-13

CAPLUS COPYRIGHT 2004 ACS on STN L11 ANSWER 1 OF 13

ACCESSION NUMBER:

2003:173639 CAPLUS

DOCUMENT NUMBER:

138:217168

TITLE:

Serine protease inhibitory glycoprotein from

shark cartilage and therapeutic uses

INVENTOR(S):

Dupont, Eric; Beliveau, Richard; Gingras, Denis;

Renaud, Alain; Cadoret, France; Dimitriadou, Violetta;

Falardeau, Pierre

PATENT ASSIGNEE(S):

Les Laboratoires Aeterna Inc., Can.

SOURCE:

PRT AB

PCT Int. Appl., 58 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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WO :	2003	01862	20		A2		2003	0306	1	VO 20	002-0	CA13	9		20	00208	323
WO :	2003	01862	20		А3		2003	0821									
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		CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KΡ,	KR,	ΚZ,	LC,	LK,	LR,
		LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NO,	NZ,	OM,	PH,
		PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,	TZ,
		UA,	UG,	US,	UZ,	VC,	VN,	ΥU,	ZA,	ZM,	ZW,	AM,	ΑZ,	BY,	KG,	KZ,	MD,
		RU,	ТJ,	TM													
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		CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	ΙE,	ΙΤ,	LU,	MC,	NL,
		PT,	SE,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,
		NE,	SN,	TD,	TG												
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mol.). The p54 glycoprotein has a protein backbone of about 46 kDa (hereinafter referred to as "p46 protein" or "p46"). The invention also relates to a process for preparing the same, methods as well as compns. for treating, preventing or alleviating the symptoms of disorders and diseases associated with an excess level of serine protease. Amongst these diseases are psoriasis, emphysema, pulmonary hypertension, liver fibrosis, anemia, diseases characterized by tumor growth or invasion, as well as any disease involving mast-cells. According to another embodiment, the present invention provides for antibodies directed specifically against p54 or p46 and methods for detecting p54 or p46 by using these specific antibodies. The activity of p54 towards other types of proteases in addition to elastases (PPE and HLE) revealed that p54 also inhibits to a lesser extent other serine proteinases such as chymotrypsin (53 %), plasmin (49 %) and trypsin (30 %), whereas it does not inhibit MMP-2 and MMP-7, cathepsin D, cathepsin G, thrombin and papain (see table III).

L11 ANSWER 2 OF 13 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2003:989927 CAPLUS

DOCUMENT NUMBER:

140:19891

TITLE:

Compositions for treatment of diseases arising from

secretion of mast cell biochemicals

Theoharides, Theoharis C. INVENTOR(S):

PATENT ASSIGNEE(S):

U.S. Pat. Appl. Publ., 8 pp., Cont.-in-part of U.S. SOURCE:

Ser. No.773,576.

CODEN: USXXCO

DOCUMENT TYPE: LANGUAGE:

Patent

FAMILY ACC. NUM. COUNT:

English

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003232100	A1	20031218	US 2003-439301	20030516
US 6689748	B1	20040210	US 1998-56707	19980408
PRIORITY APPLN. INFO.:			US 1998-56707	A3 19980408
			US 2001-773576	A2 20010202

Compns. for treatment of diseases arising from products secreted by AB activated tissue mast cells, composed of, as active ingredients, unprocessed olive kernel (pit) extract that increases absorption of these compns. in various routes of administration, and one or more of a heavily sulfated, non-bovine proteoglycan such as shark cartilage chondroitin sulfate C, a hexosamine sulfate such as D-glucosamine sulfate, a flavonoid such as quercetin, Sadenosylmethionine, a histamine-1 receptor antagonist, a histamine-3 receptor agonist, a CRH antagonist, caffeine, fragments of myelin basic protein, rutin, polyunsatd. fatty acids, Bitter Willow Extract and a polyamine.

L11 ANSWER 3 OF 13 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on

ACCESSION NUMBER: 2004:7340 BIOSIS DOCUMENT NUMBER: PREV200400008315

TITLE:

Proteoglycan compositions for treating arthritic

inflammatory conditions.

AUTHOR(S): Theoharides, Theoharis C. [Inventor, Reprint Author]

PATENT INFORMATION: US 6645482 November 11, 2003

SOURCE:

Official Gazette of the United States Patent and Trademark

Office Patents, (Nov 11 2003) Vol. 1276, No. 2. http://www.uspto.gov/web/menu/patdata.html. e-file.

ISSN: 0098-1133 (ISSN print).

DOCUMENT TYPE:

Patent English

LANGUAGE: ENTRY DATE:

Entered STN: 17 Dec 2003

Last Updated on STN: 17 Dec 2003 Compositions with synergistic anti-inflammatory effects in inflammatory AB diseases resulting from activation and consequent degranulation of mast

cells and followed by secretion of inflammatory biomolecules from the activated mast cells, composed of a heavily sulfated, non-bovine

proteoglycan such as shark cartilage chondroitin

sulfate C, and one or more of a hexosamine sulfate such as D-glucosamine sulfate, a flavone such as quercetin, an unrefined olive kernel extract that increases absorption of these compositions in various routes of administration, S-adenosylmethionine, a histamine-1 receptor antagonist, a histamine-3 receptor agonist, an antagonist of the actions of CRH, caffeine, and a polyamine.

L11 ANSWER 4 OF 13 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2004:121318 CAPLUS

DOCUMENT NUMBER:

141:179397

TITLE:

Anti-arthritic effect of a new diet-supplement

containing red ginseng extract and glucosamine complex

AUTHOR (S):

Jeong, Choon Sik; Hyun, Jin Ee; Kang, Min Hee; Sim, Joon-Soo; Son, Mi Jin; Jung, Sang Hoon; Kim, Jong

Hoon; Lee, Kwang-Seong; Kim, Yeong Shik

CORPORATE SOURCE:

College of Pharmacy, Duksung Women's University,

Seoul, 132-714, S. Korea

Saengyak Hakhoechi (2003), 34(4), 327-334 SOURCE:

> CODEN: SYHJAM; ISSN: 0253-3073 Korean Society of Pharmacognosy

PUBLISHER: DOCUMENT TYPE: Journal LANGUAGE: Korean

We evaluated the anti-arthritic effect of a new diet-supplement product containing red ginseng, glucosamine, shark cartilage,

ascorbic acid and manganese chloride for the relieving arthritic symptoms. Anti-inflammatory activities of the aqueous extract of red ginseng (250 and 500

mg/kg), glucosamine (240 mg/kg) and shark cartilage

(240 mg/kg) were tested individually on vascular permeability and

carrageenan-induced paw edema. Glucosamine and shark

cartilage showed the inhibition of vascular permeability by 29.6

and 32.9%, resp. Red ginseng (500 mg/kg) and shark

cartilage showed the inhibition of carrageenan-induced paw edema

at 0.5, 1, 2 and 3 h. The supplement (red ginseng mixture: RGM) composed of red ginseng (43.5%), glucosamine (25.0%), shark

cartilage (25.0%), ascorbic acid (5.0%) and manganese chloride (1.5%) was prepared and its inhibitory activities including vascular permeability and carrageenan-induced paw edema were comparable to anti-inflammatory drugs such as diclofenac and ibuprofen. It was also tested on adjuvant-induced arthritis in rats as one of chronic arthritic tests and Randall-Selitto assay as an analgesic test. RGM

showed the inhibition against the swelling of rat paws induced by Mycobacterium tuberculosis at a dose of 1,500 mg/kg. Determination of

cytokines

of the sera sampled from arthritis-induced animals indicated that RGM increased the levels of interferon-γ and interleukin-6, representing the immunostimulatory effect by red ginseng. RGM treatment moderately reduced the production of NO in RAW 264.7 cells in a dose-dependent manner. Taken together, these results support that RGM can be applicable for the improvement of arthritic symptoms as a new diet-supplement.

L11 ANSWER 5 OF 13 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation.

STN

ACCESSION NUMBER: 2003:578098 BIOSIS DOCUMENT NUMBER: PREV200300583728

TITLE: Clinical implications of matrix metalloproteinases.

AUTHOR(S): Mandal, Malay; Mandal, Amritlal; Das, Sudip; Chakraborti,

Tapati; Chakraborti, Sajal [Reprint Author]

CORPORATE SOURCE: Department of Biochemistry and Biophysics, University of

Kalyani, Kalyani, West Bengal, 741235, India

s chakraborti@hotmail.com

SOURCE: Molecular and Cellular Biochemistry, (October 2003) Vol.

252, No. 1-2, pp. 305-329. print.

ISSN: 0300-8177 (ISSN print).

DOCUMENT TYPE: LANGUAGE:

Article English

ENTRY DATE:

Entered STN: 10 Dec 2003

Last Updated on STN: 10 Dec 2003

Matrix metalloproteinases (MMPs) are a family of neutral proteinases that are important for normal development, wound healing, and a wide variety of pathological processes, including the spread of metastatic cancer cells, arthritic destruction of joints, atherosclerosis, pulmonary fibrosis, emphysema and neuroinflammation. In the central nervous system (CNS), MMPs have been shown to degrade components of the basal lamina, leading to disruption of the blood brain barrier and to contribute to the neuroinflammatory responses in many neurological diseases. Inhibition of MMPs have been shown to prevent progression of these diseases. Currently, certain MMP inhibitors have entered into clinical trials. A goal to the future should be to design selective synthetic inhibitors of MMPs that have minimum side effects. MMP inhibitors are designed in such a way that these can not only bind at the active site of the proteinases but also to

have the characteristics to bind to other sites of MMPs which might be a promising route for therapy. To name a few: catechins, a component isolated from green tea; and Novastal, derived from extracts of shark cartilage are currently in clinical trials for the treatment of MMP-mediated diseases.

L11 ANSWER 6 OF 13 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:594640 CAPLUS

DOCUMENT NUMBER:

137:145588

TITLE:

Proteoglycan compositions for treatment of

inflammatory conditions Theoharides, Theoharis C.

INVENTOR(S):

PATENT ASSIGNEE(S):

SOURCE: PCT Int. Appl., 21 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

		rent 1								AP		ATION				ATE	
										WO						0020	 103
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		2002															
								US									
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	EP	1365	777			A2		2003	1203	EP	2002	2-7053	707		2	0020	103
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			ΙE,	FI,	CY,	TR										·	•
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	US	6645	482			B2		2003	1111								
		2003		88		<b>A</b> 1		2003	0605	US	2002	-3293	367		2	0021	227
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		66418						2003	1104								
		20040						2004	0108			-6109				0030'	702
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Compns. with synergistic anti-inflammatory effects in inflammatory AB diseases resulting form activation and consequent degranulation of mast cells and followed by secretion of inflammator biomols. from the activated mast cells, composed of a heavily sulfated, non-bovine proteoglycan such as shark cartilage chondroitin sulfate C, and one or more of a hexosamine sulfate such as D-glucosamine sulfate, a flavone such as quercetin, an unrefined kernel olive oil that increases absorption of these compns. in various routes of administration, S-adenosylmethionine, a histamine-1 receptor antagonist, a histamine-3 receptor agonist, an antagonist of the actions of CRH, caffeine, and a polyamine. For example, a composition for protecting against cardiovascular disease, in the form of capsule to be taken 2 capsules orally 2-3 times per day, contained chondroitin sulfate 50 mg, kaempferol 100 mg, S-adenosylmethionine 50 mg, niacin 100 mg, and unrefined kernel olive oil 900-1200 mg.

L11 ANSWER 7 OF 13 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:960660 CAPLUS

DOCUMENT NUMBER:

138:19488

TITLE:

Method and pharmaceutical compositions using anti-microtubule agents for treating multiple sclerosis and other inflammatory diseases

Hunter, William L.

PATENT ASSIGNEE(S): Angiotech Pharmaceuticals, Inc., Can.

SOURCE: U.S., 180 pp., Cont.-in-part of U.S. Appl. 2002

37,919.

CODEN: USXXAM

DOCUMENT TYPE:

LANGUAGE:

INVENTOR(S):

Patent English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

	TENT				KIN		DATE				ICAT					ATE	
	6495		10		B1		2002				.998-					9980	
	2002		19		A1		2002			US 1	997-	9805	49		1	9971	201
	6515				B2		2003										
	1070				A2		2001			EP 2	000-	1235	57		1	9971	202
	1070				A3		2001										
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	R:			CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
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EP	1090	637			A2		2001	0411		EP 2	000-	1235	37		1	9971	202
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	R:	ΑT,	BE,	CH,	DΕ,	DK,	ES,	FR,	GB,	GR,	ΙT,	LI,	LU,	NL,	SE,	MC,	PT,
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EP	1092	433			B1		2003	0806									
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
		ΙE,	FI										•	•	•	•	•
JP	2002	2263	99		A2		2002	0814		JP 2	001-	4018	99		1	9971:	202
WO	9962	510			A2		1999	1209			999-0					9990	
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		KG,	KP,	KR.	KZ.	LC.	LK,	LR.	LS.	LT.	LU.	LV.	MD.	MG.	MK.	MNI	MW
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AB Methods and compns. for treating or preventing inflammatory diseases, e.g. psoriasis or multiple sclerosis, are provided, comprising delivering to the site of inflammation an anti-microtubule agent (e.g. paclitaxel), or analog or derivative thereof.

REFERENCE COUNT:

THERE ARE 171 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE REFORMAT

L11 ANSWER 8 OF 13 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2001:63838 CAPLUS 134:120960

DOCUMENT NUMBER: TITLE:

Green-lip mussel product compositions addressing

inflammation and/or degenerative disorders INVENTOR(S): Hashmi, Syed Ziauddin; Leech, Wayne Frederick;

Mclaren, Donald George; McSporran, Keith David

PATENT ASSIGNEE(S): Bomac Laboratories Limited, N. Z.

SOURCE:

PCT Int. Appl., 30 pp. CODEN: PIXXD2

DOCUMENT TYPE:

LANGUAGE .

Patent English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO. KIND DATEAPPLICATION NO. DATE -----\_ \_ \_ \_ ----------WO 2001005411 A1 20010125 WO 2000-NZ135 20000721 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG NZ 336856 Α 20010330 NZ 1999-336856 19990721 AU 727355 В3 20001214 AU 2000-48775 20000721 JP 2003504408 T2 20030204 JP 2001-510466 20000721 AU 761829 B2 20030612 AU 2000-63257 20000721 EP 1408999 A1 20040421 EP 2000-950109 20000721 AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY PRIORITY APPLN. INFO.: NZ 1999-336856 A 19990721 NZ 1999-500630 A 19991027 NZ 2000-505875 A 20000721

W 20000721 The present invention is directed to compns. for primarily addressing AB degenerative complaints, in particular joint related conditions, such as arthritis and rheumatism, in which there may also be associated inflammation. Other potential uses are also discussed, as well as prophylactic and curative applications. Preferred embodiments incorporate green-lip mussel products (particularly GLME) with shark cartilage or chondroitin compds. Plant and bark based antioxidants are employed in a number of embodiments. Dosage forms containing GLME, shark cartilage, vitamins and minerals are given. The compns. can be used especially for animals.

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 9 OF 13 MEDLINE on STN DUPLICATE 1

ACCESSION NUMBER:

2001131805 MEDLINE

WO 2000-NZ135

DOCUMENT NUMBER:

PubMed ID: 11196523

TITLE:

Surfing the Net--information on the World Wide Web for

persons with arthritis: patient empowerment or

patient deceit?.

COMMENT:

Comment in: J Rheumatol. 2001 Jan; 28(1):1-2. PubMed ID:

11196508

AUTHOR:

Suarez-Almazor M E; Kendall C J; Dorgan M

CORPORATE SOURCE:

Veteran Affairs Medical Center, Health Services, Research, Baylor College of Medicine, Houston, Texas 77030, USA..

mes@bcm.tmc.edu

SOURCE:

Journal of rheumatology, (2001 Jan) 28 (1) 185-91.

Journal code: 7501984. ISSN: 0315-162X.

PUB. COUNTRY:

Canada

DOCUMENT TYPE:

(EVALUATION STUDIES)

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE:

English

FILE SEGMENT:

Priority Journals

ENTRY MONTH:

200103

ENTRY DATE:

Entered STN: 20010404

Last Updated on STN: 20010404 Entered Medline: 20010301

AB OBJECTIVE: In the past few years access to the Internet has become readily available. Patients are increasingly seeking and obtaining health information through the Internet, most often the World Wide Web (WWW). assessed the content, authorship, and scope of the information available on WWW in relation to rheumatoid arthritis. METHODS: In an attempt to replicate use by the average person, a broad search of the Internet was conducted for the phrase "rheumatoid arthritis" using WebCrawler, a commonly used search engine. All the "hits" were critically assessed after visiting and collecting information from the respective Web sites in relation to relevance, scope, authorship, type of publication, and financial objectives. RESULTS: The search returned 537 hits. We evaluated 531-2 did not exist, 2 could not be contacted, one was not in English, and one required a membership to access. The 531 hits originated from 388 Web sites. Only 198 (51%) were considered to be relevant and 7 (2%) were of doubtful relevance. Thirty-four (17%) were posted by an individual, 57 (28%) by a nonprofit organization, 104 (51%) by a profit industry, and 10 (5%) by universities. Ninety-one (44%) promoted alternative therapies, the most common including cetyl-myristoleate, colloidal minerals, Pycnogenol, shark cartilage, and Tahitian Noni. Of the 107 sites with financial interests, 76 (71%) promoted alternative medicine. The first 100 hits only identified about a third of the nonprofit organizations or university owned Web pages. CONCLUSION: Many sites easily accessed by consumers appear to be profit based companies advertising an alternative product claimed to be effective for many conditions. These findings emphasize the need for critical evaluation of Web site contents.

L11 ANSWER 10 OF 13 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1999:731749 CAPLUS

DOCUMENT NUMBER:

131:332105

TITLE:

Inhibition of angiogenesis by sea cucumber fractions

containing sulfated polysaccharides

INVENTOR (S):

Collin, Peter Donald

PATENT ASSIGNEE(S):

Coastside Bio Resources, USA

SOURCE:

U.S., 9 pp., Cont.-in-part of U.S. Ser. No. 692,175,

abandoned. CODEN: USXXAM

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5985330	Α	19991116	US 1997-880359	19970623
ORITY APPLN. INFO.:			US 1996-692175	19960805

PRTO The present invention provides inhibition of angiogenesis in a warm-blooded animal by the administration of prepns. isolated from the echinoderm sea cucumber (Class Holothuroidea). The preparation contains sulfated polysaccharides, sterol glycosides, saponins, lactones, peptides, protamines, glycogens, saccharides, and polysaccharides, and is useful as a therapeutic agent against malignant tumors and as a preventive or therapeutic drug against various diseases, such as rheumatoid arthritis, caused by vascular hyperplasia. Fucosylated chondroitin sulfate was extracted from the body wall of the sea cucumber Ludwigothurea grisea by papain digestion and tested for anti-angiogenic activity using the Chorioallantoic Membrane Assay (CAM) method. Sea cucumber fucosylated chondroitin sulfate showed good anti-angiogenic activity. The activity seen was nearly as high as that of the pos. control, hydrocortisone/heparin, and higher than that seen with

shark cartilage chondroitin-6-sulfate. A fraction termed B1000, consisting of sea cucumber epithelium, at 100  $\mu g/mL$ decreased the ability of human melanoma tumor cell line C8161 to invade by the value of 63% inhibition.

REFERENCE COUNT:

THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS 16 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 11 OF 13 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1995:958356 CAPLUS

DATE

DOCUMENT NUMBER:

123:350289

TITLE:

Dietary supplement for pain relief

INVENTOR (S): Woodward, Robert John

PATENT ASSIGNEE(S):

SOURCE:

Brit. UK Pat. Appl., 9 pp.

CODEN: BAXXDU

DOCUMENT TYPE:

Patent

LANGUAGE:

English

KIND

FAMILY ACC. NUM. COUNT:

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PATENT INFORMATION:

PATENT NO.

GB 2286528	A1	19950823	GB 1994-3063	` 19940217
GB 2286528	B2	19980916		13310217
PRIORITY APPLN. INFO.:			GB 1994-3063 `	19940217
AB A dietary supplement	contai	ning sources	of vitamin's B3,	B5, and/or B6.
D-phenylalanine, glu	cosamin	ne sulfate, ar	nd optionally muc	copolysaccharides
such as chondroitin	sulfate	and shark ca	artilage can	
provide relief of jo	int or	muscular pair	n, e.g. arthriti:	s. Thus,

a tablet formulation contained pantothenic acid 100, shark cartilage 100, DL-phenylalanine 50, chondroitin sulfate 50, glucosamine sulfate 50 mg, and conventional tableting additives.

L11 ANSWER 12 OF 13 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1993:623380 CAPLUS

DOCUMENT NUMBER:

119:223380

TITLE:

The clinical significance on keratan sulfate levels of serum and synovial fluid in cases of osteoarthritis of

APPLICATION NO.

DATE

the knee

AUTHOR (S):

Mibe, Junya

CORPORATE SOURCE: SOURCE:

Dep. Orthopedic Surg., Tokyo Med. Coll., Tokyo, Japan

Tokyo Ika Daigaku Zasshi (1993), 51(2), 141-52

CODEN: TIDZAH; ISSN: 0040-8905

DOCUMENT TYPE:

Journal

LANGUAGE:

Japanese

AB Keratan sulfate (KS) is a sulfated glycosaminoglycan mostly found in the extracellular matrix of cartilage and is thought to be a useful marker of cartilage metabolism The purpose of this study was to measure serum and synovial fluid KS levels and to clarify its clin. significance. Serum samples were obtained from 311 healthy adults and 93 cases of primary osteoarthritis (OA) of the knee. OA cases were subdivided into 3 groups; mild OA group, moderate OA group, and severe OA group. Specimens of synovial fluid of the knee were obtained from 79 OA cases (subdivided into the same 3 groups as above), 7 cases of acute traumatic synovitis (TS), and 22 cases of fresh medial collateral ligament injury (LI) of the knee. A modification of the competitive ELISA method of Thonar was employed, and the standard antigen was keratan polysulfate from shark cartilage. The normal serum KS level was 239.4 ± 55.7 ng/mL. There was no difference according to sex, but there was correlation with age (r = 0.35). The serum KS level of OA cases was  $251.4 \pm 64.5$  ng/mL. There was no difference from normal subjects, or among each grade. The synovial fluid KS level of OA was 8.39  $\pm$  7.78  $\mu$ g/mL. There were differences among each grade; the mild OA group was 11.37 ± 9.34  $\mu g/mL,$  the moderate OA group was 6.38  $\pm$  3.60  $\mu g/mL,$  and the severe OA group was 2.87  $\pm$  1.21  $\mu$ g/mL. When the mild OA group was

subdivided into acute stage and chronic stage, the KS level of the acute stage was higher than that of the chronic stage. Furthermore, the synovial fluid KS level of TS cases was 18.85  $\pm$  9.76  $\mu g/mL$  while that of LI cases was 94.43  $\pm$  87.42  $\mu g/mL$ , therefore both cases were higher than the OA cases. It was concluded that the serum KS level reflects general factors. However, several cases which have high KS levels may suggest some abnormality in local cartilage metabolic activity. The synovial fluid KS level is associated with the grade of OA or severity of injury, reflecting local cartilage metabolism, especially catabolism.

L11 ANSWER 13 OF 13 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1981:145330 CAPLUS

DOCUMENT NUMBER:

94:145330

TITLE:

Cartilage extraction processes and products

INVENTOR(S):

Balassa, Leslie L. Lescarden Ltd., USA

PATENT ASSIGNEE(S): SOURCE:

PCT Int. Appl., 34 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 8002501	A1	19801127	WO 1980-US503	19800502
W: JP, RO, SU				
RW: AT, CH, DE,	FR, GB	, LU, NL, SE		
US 4350682	Α	19820921	US 1980-137547	19800404
EP 28254	A1	19810513	EP 1980-901098	19800502
EP 28254	B1	19840801		-
R: AT, CH, DE,	FR, GB	, LU, NL, SE		
AT 8738	E	19840815	AT 1980-901098	19800502
CA 1140851	A1	19830208	CA 1980-351614	19800509
PRIORITY APPLN. INFO.:			US 1979-38051	19790511
			US 1980-137547	19800404
•			EP 1980-901098	19800502
			WO 1980-US503	19800502

AB Cartilage exts. suitable for use in cosmetics, pharmaceuticals, and foods were prepared from bovine trachea, shark spinal column, or other animal source. Thus, 5 kg beef trachea and 50 g H2O were cooked, in the absence of air, at 20 psig for 4 h with stirring at 20-60 rpm in a steam-jacketed vessel, the contents were filtered at 90°, and the fibrous matter was pressed (5 psi). The fluids were centifuged to give 2500 g containing 840 g fat and 1550 g protein matter. The protein portion gelled on refrigeration, had a taste similar to meat extract, and could be used as a dietary supplement or as a pharmaceutical preparation

---Logging off of STN---

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